

## PATENT COOPERATION TREATY

## PCT

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## INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

(Chapter II of the Patent Cooperation Treaty)

(PCT Article 36 and Rule 70)

Applicant's or agent's file reference PV/448/PCT	<b>FOR FURTHER ACTION</b>		See Form PCT/PEA/416
International application No. PCT/CZ2005/000010	International filing date (day/month/year) 03.02.2005	Priority date (day/month/year) 05.02.2004	
International Patent Classification (IPC) or national classification and IPC INV. C07C303/40			
Applicant ZENTIVA, A.S.			
<p>1. This report is the international preliminary examination report, established by this International Preliminary Examining Authority under Article 35 and transmitted to the applicant according to Article 36.</p> <p>2. This REPORT consists of a total of 5 sheets, including this cover sheet.</p> <p>3. This report is also accompanied by ANNEXES, comprising:</p> <p>a. <input checked="" type="checkbox"/> sent to the applicant and to the International Bureau a total of 5 sheets, as follows:</p> <p><input checked="" type="checkbox"/> sheets of the description, claims and/or drawings which have been amended and are the basis of this report and/or sheets containing rectifications authorized by this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions).</p> <p><input type="checkbox"/> sheets which supersede earlier sheets, but which this Authority considers contain an amendment that goes beyond the disclosure in the international application as filed, as indicated in item 4 of Box No. I and the Supplemental Box.</p> <p>b. <input type="checkbox"/> (sent to the International Bureau only) a total of (indicate type and number of electronic carrier(s)) , containing a sequence listing and/or tables related thereto, in electronic form only, as indicated in the Supplemental Box Relating to Sequence Listing (see Section 802 of the Administrative Instructions).</p>			
<p>4. This report contains indications relating to the following items:</p> <p><input checked="" type="checkbox"/> Box No. I Basis of the report</p> <p><input type="checkbox"/> Box No. II Priority</p> <p><input type="checkbox"/> Box No. III Non-establishment of opinion with regard to novelty, inventive step and industrial applicability</p> <p><input type="checkbox"/> Box No. IV Lack of unity of invention</p> <p><input checked="" type="checkbox"/> Box No. V Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement</p> <p><input type="checkbox"/> Box No. VI Certain documents cited</p> <p><input type="checkbox"/> Box No. VII Certain defects in the international application</p> <p><input checked="" type="checkbox"/> Box No. VIII Certain observations on the international application</p>			
Date of submission of the demand  01.12.2005		Date of completion of this report  20.04.2006	
Name and mailing address of the international preliminary examining authority:  European Patent Office D-80298 Munich Tel. +49 89 2399 - 0 Tx: 523656 epmu d Fax: +49 89 2399 - 4465		Authorized officer  Grammenoudi, S  Telephone No. +49 89 2399-8324	



**INTERNATIONAL PRELIMINARY REPORT  
ON PATENTABILITY**

International application No.  
PCT/CZ2005/000010

**Box No. I Basis of the report**

1. With regard to the **language**, this report is based on the international application in the language in which it was filed, unless otherwise indicated under this item.
- ☐ This report is based on translations from the original language into the following language , which is the language of a translation furnished for the purposes of:
- ☐ international search (under Rules 12.3 and 23.1(b))
  - ☐ publication of the international application (under Rule 12.4)
  - ☐ international preliminary examination (under Rules 55.2 and/or 55.3)
2. With regard to the **elements\*** of the international application, this report is based on *(replacement sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to this report)*:

**Description, Pages**

1, 2, 4-8, 10-14	as originally filed
3, 9	received on 06.12.2005 with letter of 01.12.2005

**Claims, Numbers**

1-10	received on 06.12.2005 with letter of 01.12.2005
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- ☐ a sequence listing and/or any related table(s) - see Supplemental Box Relating to Sequence Listing

3. ☐ The amendments have resulted in the cancellation of:
- ☐ the description, pages
  - ☐ the claims, Nos.
  - ☐ the drawings, sheets/figs
  - ☐ the sequence listing (*specify*):
  - ☐ any table(s) related to sequence listing (*specify*):
4. ☐ This report has been established as if (some of) the amendments annexed to this report and listed below had not been made, since they have been considered to go beyond the disclosure as filed, as indicated in the Supplemental Box (Rule 70.2(c)).
- ☐ the description, pages
  - ☐ the claims, Nos.
  - ☐ the drawings, sheets/figs
  - ☐ the sequence listing (*specify*):
  - ☐ any table(s) related to sequence listing (*specify*):

\* If item 4 applies, some or all of these sheets may be marked "superseded."

**INTERNATIONAL PRELIMINARY REPORT  
ON PATENTABILITY**

International application No.  
PCT/CZ2005/000010

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**Box No. V Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement**

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1. Statement

Novelty (N)	Yes: Claims	1-10
	No: Claims	
Inventive step (IS)	Yes: Claims	1-10
	No: Claims	
Industrial applicability (IA)	Yes: Claims	1-10
	No: Claims	

2. Citations and explanations (Rule 70.7):

**see separate sheet**

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**Box No. VIII Certain observations on the international application**

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The following observations on the clarity of the claims, description, and drawings or on the question whether the claims are fully supported by the description, are made:

**see separate sheet**

D1= EP-A-0 257 787

1. The present application relates to a process for producing (R)-(-)-5-(2-aminopropyl)-2-methoxybenzenesulfonamide.
2. Document D1 cited on page 3, line 2, page 4, line 28 and page 10, lines 1 and 11 is considered to represent the closest state of the art. It discloses a process for producing (R)-(-)-5-(2-aminopropyl)-2-methoxybenzenesulfonamide which comprises reacting 5-acetyl-2-methoxybenzenesulfonamide with R-(+)- $\alpha$ -methylbenzylamine to obtain 2R,1R-2-methoxy-5-[2-(1-methylbenzylamino)propyl]benzene-sulfonamide hydrochloride, hydrogenating the latter to (R)-(-)-5-(2-aminopropyl)-2-methoxybenzenesulfonamide hydrochloride whereupon this salt is converted to (R)-(-)-5-(2-aminopropyl)-2-methoxybenzenesulfonamide (see D1, Examples 1-3). The problem to be solved by the present application with respect to the cited prior art is to provide an alternative process for the same purpose.
3. The process according to present claim 1 differs from the known method in that it starts with the amine of formula VIII which is subsequently N-acylated, chlorosulfonated, hydrogenated and deacylated to produce (R)-(-)-5-(2-aminopropyl)-2-methoxybenzenesulfonamide. There is no suggestion of such a process in the state of the art.

Accordingly, the subject-matter of claim 1 meets the requirements of Article 33(2) and (3) PCT.

4. The compounds of claim 9 are novel and serve as intermediates in the process according to claim 1. Thus, the subject-matter of claim 9 meets the requirements of Article 33(2) and (3) PCT.
5. Dependent claims 1-8 and 10 concern particular embodiments of claims 1 and 9 respectively. They fulfil the requirements of Art. 33(2) and (3) PCT as well.

## SECTION VIII

**INTERNATIONAL PRELIMINARY  
REPORT ON PATENTABILITY  
(SEPARATE SHEET)**

International application No.

PCT/CZ2005/000010

1. The feature "50-100°C" of claim 3 is not supported by the description (cf. page 5, lines 11-12) as required by Article 6 PCT.
2. The number "VII" on page 9, line 4 and page 12, line 6 as well as the formula of R-(+)- $\alpha$ -methylbenzylamine depicted on page 10, first reaction scheme (cf. Example 6, 2.1) are obviously incorrect (Art. 6 PCT).
3. The two-step chlorosulfonation reaction scheme on page 9 as originally filed has been replaced by the hydrogenation reaction scheme depicted on page 10, lines 1-5 (with a corrected formula of R-(+)- $\alpha$ -methylbenzylamine). As a result, the description of the prior art method for producing tamsulosin now comprises two hydrogenations on platinum wherein, however, the compound numbering (VII and III) as well as the yield data (40%, 74% and 30%) of the first hydrogenation step from the chlorosulfonation. This inconsistency within Example 6 renders it unclear, contrary to the requirements of Article 6 PCT.

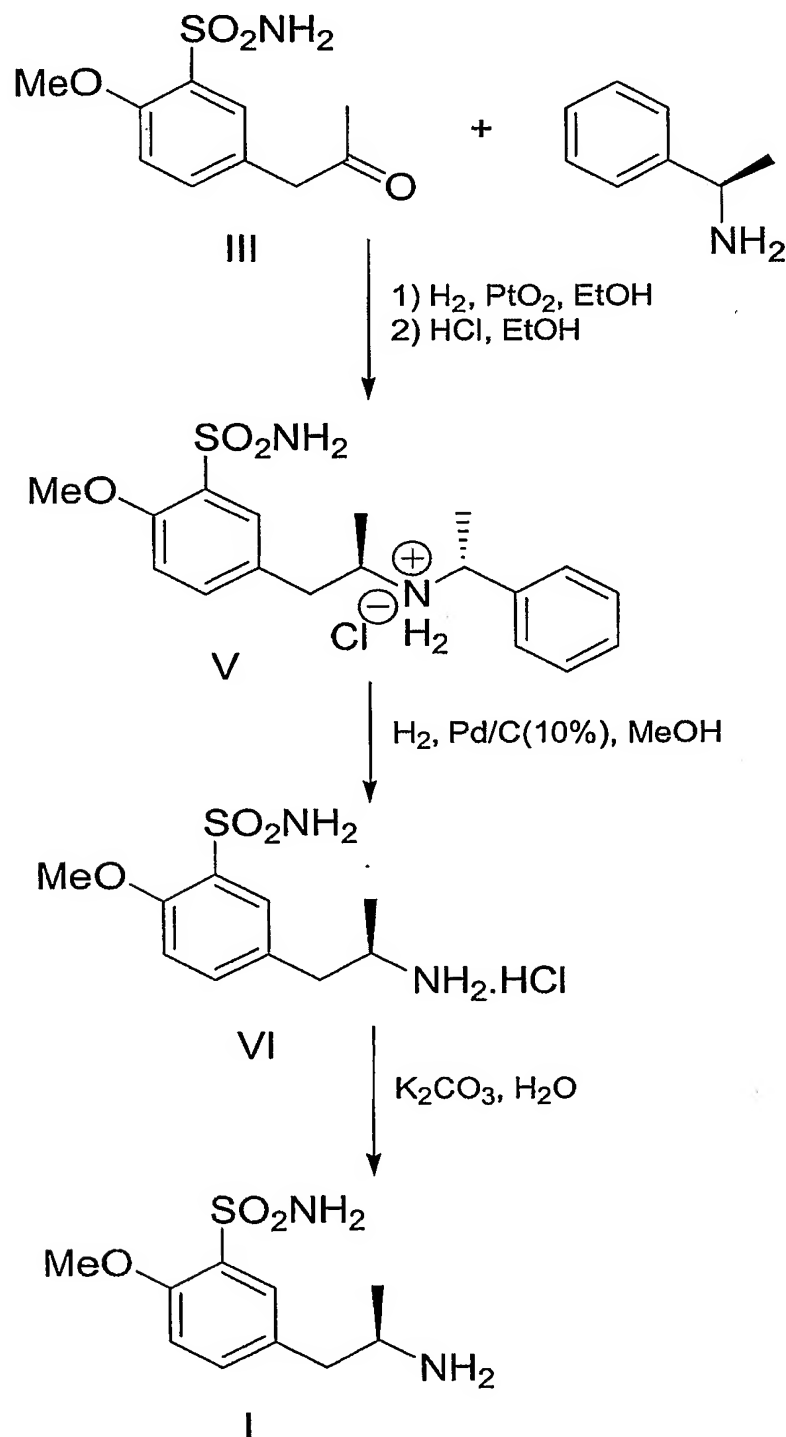
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IV

The synthesis turned out to be very advantageous if the starting amine I was prepared according to the method described in patent EP 257787, or divisional EP 380144.

5

The patents describe the path indicated in the following scheme:



AMENDED SHEET

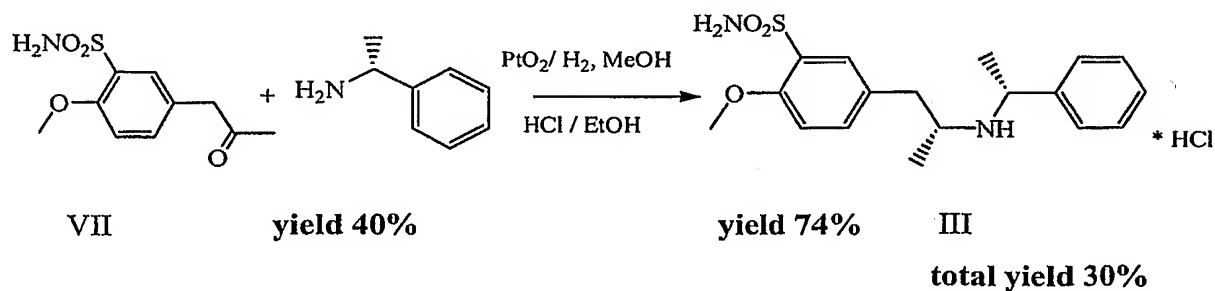
#### Preparation of (R)-(-)-5-(2-aminopropyl)-2-methoxybenzenesulfonamide I

<sup>1</sup>H NMR: δ 0.99(d, J=6.2, 3H, CH<sub>3</sub>); 2.54(dd, J=13.6 J=6,8, 1H, CH<sub>2</sub>); 2.59 (dd, J=13.6; J=6.7; 1H, CH<sub>2</sub>); 3.00(sex, J=6.4; 1H, CH); 3.53(bs, NH<sub>2</sub>); 3.92(s, 3H, CH<sub>3</sub>O); 7.16(d, J=8.4; 1H, CH<sub>arom</sub>); 7.41(dd, J=8.4; J= 2.2; 1H, CH<sub>arom</sub>); 7.59(d, J=2.2; 1H, CH<sub>arom</sub>). (CD<sub>3</sub>SOCD<sub>3</sub>, 30 °C)

**A method of preparation of tamsulosin (II) according to the prior art in comparison with the method with included steps according to the invention**

### 1.1. Chlorosulfonation in two steps according to US 4,731,478 (1988) :

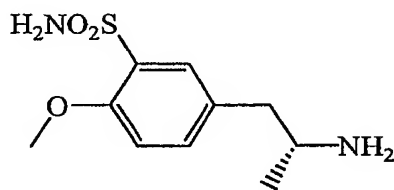
1.2.



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## C L A I M S

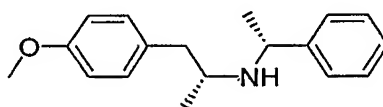
1. A method of preparation of (R)-(-)-5-(2-aminopropyl)-2-methoxybenzenesulfonamide  
of formula I



I

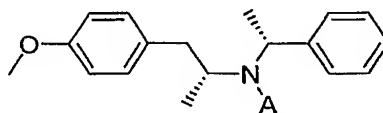
characterized in that

- a. a protecting group is introduced to N-[(1R)-2-(4-methoxyphenyl)-1-methylethyl]-N-[(1R)-1-phenylethyl]amine of formula VIII



VIII

to obtain an amide of formula IX



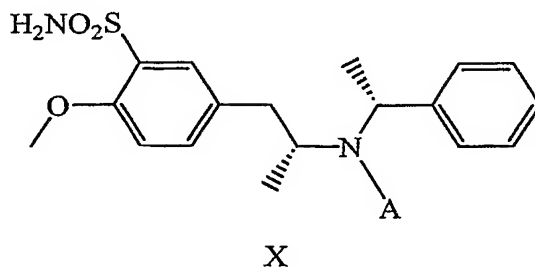
IX,

wherein A can be an acyl having 2 to 8 carbons,

- b. whereupon the amide of formula IX is chlorosulfonated and the resulting sulfochloride is converted to a sulfonamide of formula X



16



wherein A is as defined above,

5

c. and the sulfonamide of formula X is hydrogenated and deacylated to obtain the compound of formula I.

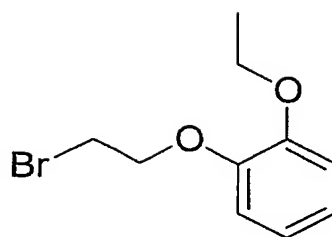
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2. The method according to claim 1 characterized in that the protecting group A is an acyl, preferably acetyl.
3. The method according to claim 2 characterized in that acetic anhydride at 50 to 100 °C is used as the acetylation agent.
4. The method according to claim 1 characterized in that the sulfochloride resulting from chlorosulfonation is not isolated and is directly converted to the sulfonamide with ammonia.
5. The method according to claim 4 characterized in that chlorosulfonation takes place in methylenechloride at -30 to +30 °C.
6. The method according to claim 1 characterized in that hydrogenation is carried out under catalysis with palladium.
7. The method according to claim 6 characterized in that the catalyst is 3% Pd/C with 50% water content at a pressure of 2 MPa and a temperature of 80 to 85 °C.
8. The method according to claim 1, characterized in that the product of formula I is further reacted with a compound of formula IV

15

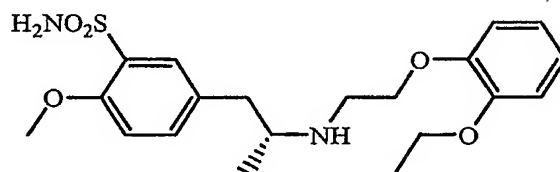
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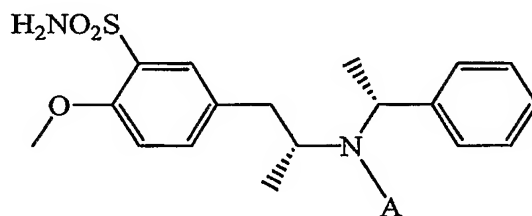
IV

to form the final product of formula II



II

9. A sulfonamide of formula X



X

wherein A is as defined in claim 1.

10. The sulfonamide according to claim 9, wherein A is acetyl.